EXHIBIT 1

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IN THE UNITED STATES DISTRICT COURT FOR THE DISTRICT OF MASSACHUSETTS

IN RE:

) CA No. 01-12257-PBS
PHARMACEUTICAL INDUSTRY AVERAGE
WHOLESALE PRICE LITIGATION
) Pages 1 - 87

SETTLEMENT HEARING

BEFORE THE HONORABLE PATTI B. SARIS UNITED STATES DISTRICT JUDGE

United States District Court 1 Courthouse Way, Courtroom 19 Boston, Massachusetts June 13, 2011, 2:20 p.m.

LEE A. MARZILLI
OFFICIAL COURT REPORTER
United States District Court
1 Courthouse Way, Room 7200
Boston, MA 02210
(617)345-6787

Page 7 1 reasons for that are: We didn't have a trial verdict or trial 2 benchmarks like we did with AstraZeneca or BMS, and we had to 3 evaluate the liability case, and we just didn't have the hot documents, you know, showing pushing the spread to physicians 5 like we did with AstraZeneca or with BMS. So the liability 6 case wasn't nearly as strong, and we feel that the settlement 7 of \$100 million is substantial, given those risks. Well, wasn't that originally how we 9 decided fast track versus non-fast track, Track One, Track Two? 10 Is that the reason we triaged initially? You cherry-picked the 11 other five. 12 MR. BERMAN: Right, we cherry-picked some 13 multi-source, some not multi-source, yes, so that we put our 14 best case in fast track, AstraZeneca. 15 With respect to the multi-source drug, Dr. Hartman 16 calculated \$800 million of damages, but that was using the 17 30 percent yardstick; and the damages, for the reasons I'll 18 explain in a moment, are probably nonexistent for Class B. 19 First of all, we have the J-Code issue, and on the third page 20 of our slides I had quoted from your trial verdict in which you 21 identified the problem that we would have is, we can't match up 22 any single patient's use of multi-source drug with the 23 defendant manufacturer. So we faced a real problem in that 24 regard. And we were going to propose some market share theory,

but, you know, those theories have not been warmly received by

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